

## Special Issue

# Molecular Aspects of Retinopathy and Protection

### Message from the Guest Editor

Molecular retinopathy studies encompass a wide range of knowledge that includes genetic mutations or biochemical functions that affect the retina's normal functioning.

Protection against molecular retinopathy involves a multi-faceted approach, incorporating early diagnosis, genetic counseling, lifestyle modifications, and emerging therapies. Genetic screening and molecular diagnostics play a crucial role in identifying at-risk individuals and enabling early intervention.

Innovative treatments, including gene therapy, stem cell therapy, and the use of neuroprotective agents, are being explored to address the underlying molecular causes of these retinopathies. The goal of these therapies is not only to halt disease progression but also to restore vision and improve the quality of life for affected individuals.

In summary, molecular retinopathy represents a complex interplay of genetic and biochemical factors leading to retinal degeneration. Advances in understanding the molecular underpinnings of these conditions have paved the way for novel protective and therapeutic strategies aimed at preserving vision and enhancing retinal health.

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### Deadline for manuscript submissions

closed (25 November 2024)



## International Journal of Molecular Sciences

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Impact Factor 4.9  
CiteScore 9.0  
Indexed in PubMed



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*International Journal of  
Molecular Sciences*  
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